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<u>L11</u>	adenovir\$ with (bmp or egr or hsp70 or hsp27 or hsp40 or hsp60)	38	<u>L11</u>
<u>L10</u>	L9 and 14	11	<u>L10</u>
<u>L9</u>	stress near3 (induce\$ or relat\$)	28453	<u>L9</u>
<u>L8</u>	(adenovir\$ near5 replication-deficien\$) with (bmp or egr or hsp70 or hsp27 or hsp40 or hsp60)	0	<u>L8</u>
<u>L7</u>	(adenovir\$ near5 replication-deficien\$) with (bmp or egr or wound adj heal\$)	0	<u>L7</u>
<u>L6</u>	(adenovir\$ near5 replication-deficien\$) with bmp or egr or wound adj heal\$	15073	<u>L6</u>
<u>L5</u>	13 and 14	63	<u>L5</u>
<u>L4</u>	adenovir\$ near5 replication-deficien\$	356	<u>L4</u>
<u>L3</u>	11 or L2	17932	<u>L3</u>
<u>L2</u>	bmp or bone adj morphogenic adk (protein or polypeptide)	3631	<u>L2</u>
<u>L1</u>	wound adj heal\$ or egr	15073	<u>L1</u>

END OF SEARCH HISTORY

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FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 16:34:32 ON 28 APR 2003

- L1 325115 S STRESS(3A) (RELAT? OR INDUC?) OR HYPOXIA
- L2 366 S ADENOVIR?(S) L1
- L3 120 S ADENOVIR? (10A) L1
- L4 65 DUP REM L3 (55 DUPLICATES REMOVED)
- L5 0 S L4@PD<19951231
- L6 0 S L4@PD<1996
- L7 0 S @PD<19961231 AND L4
- => d au ti so 30-65 14
- L4 ANSWER 30 OF 65 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 15
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(FILE 'HOME' ENTERED AT 15:25:30 ON 28 APR 2003)

FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 15:25:41 ON 28 APR 2003

- L1 97859 S WOUND (3A) HEAL? OR EGR
- L2 2337 S ADENOVIR? (5A) REPLICATION-DEFICIENT
- L3 32 S L1 AND L2
 - 14 DUP REM L3 (18 DUPLICATES REMOVED)
- L5 17729 S ADENOVIR? (W) VECTOR
- L6 137 S L1 AND L5
- L7 84 DUP REM L6 (53 DUPLICATES REMOVED)
- L8 75 S L1(S)L5
- L9 37 DUP REM L8 (38 DUPLICATES REMOVED)
- L10 14638 S BMP OR BONE (W) MORPHOGENIC (W) (PROTEIN OR POLYPEPTIDE)
- L11 132 S L10 AND L5
- L12 10 S L2 AND L10
- L13 91 S L10(S)L5
- L14 36 DUP REM L13 (55 DUPLICATES REMOVED)
- L15 3 DUP REM L12 (7 DUPLICATES REMOVED)

=> d au ti so 1-3 115

L15 ANSWER 1 OF 3 MEDLINE

DUPLICATE 1

- AU Okubo Y; Bessho K; Fujimura K; Iizuka T; Miyatake S I
- TI In vitro and in vivo studies of a bone morphogenetic protein-2 expressing adenoviral vector.
- SO JOURNAL OF BONE AND JOINT SURGERY. AMERICAN VOLUME, (2001) 83-A Suppl 1 (Pt 2) S99-104.

 Journal code: 0014030. ISSN: 0021-9355.
- L15 ANSWER 2 OF 3 MEDLINE

DUPLICATE 2

- AU Haberberger T C; Kupfer K; Murphy J E
- TI Profiling of genes which are differentially expressed in mouse liver in response to adenoviral vectors and delivered genes.
- SO GENE THERAPY, (2000 Jun) 7 (11) 903-9. Journal code: 9421525. ISSN: 0969-7128.
- L15 ANSWER 3 OF 3 MEDLINE

DUPLICATE 3

- AU Okubo Y; Bessho K; Fujimura K; Iizuka T; Miyatake S
- TI Expression of bone morphogenetic protein-2 via adenoviral vector in C2C12 myoblasts induces differentiation into the osteoblast lineage.
- SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (1999 Sep 7) 262 (3) 739-43.
 - Journal code: 0372516. ISSN: 0006-291X.

=> d au ti so 1-36 l14

L14 ANSWER 1 OF 36 MEDLINE

DUPLICATE 1

- AU Kaihara Shinji; Bessho Kazuhisa; Okubo Yasunori; Sonobe Junya; Komatsu Yasato; Miura Masako; Miyatake Shin-Ichi; Nakao Kazuwa; Iizuka Tadahiko
- TI Over expression of bone morphogenetic protein-3b (BMP-3b) using an adenoviral vector promote the osteoblastic differentiation in C2C12 cells and augment the bone formation induced by bone morphogenetic protein-2 (BMP-2) in rats.
- bone morphogenetic protein-2 (BMP-2) in rats.

 SO LIFE SCIENCES, (2003 Feb 28) 72 (15) 1683-93.

 Journal code: 0375521. ISSN: 0024-3205.
- L14 ANSWER 2 OF 36 MEDLINE

- AU Nishihara Ayako; Fujii Makiko; Sampath T Kuber; Miyazono Kohei; Reddi A Hari
- TI Bone morphogenetic protein signaling in articular chondrocyte

differentiation.

- SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (2003 Feb 7) 301 (2) 617-22.

 Journal code: 0372516. ISSN: 0006-291X.
- L14 ANSWER 3 OF 36 MEDLINE DUPLICATE 3
- AU Gelse Kolja; von der Mark Klaus; Aigner Thomas; Park Jung; Schneider Holm
- TI Articular cartilage repair by gene therapy using growth factor-producing mesenchymal cells.
- SO ARTHRITIS AND RHEUMATISM, (2003 Feb) 48 (2) 430-41. Journal code: 0370605. ISSN: 0004-3591.
- L14 ANSWER 4 OF 36 CAPLUS COPYRIGHT 2003 ACS
- AU Palmer, Glyn D.; Gouze, Elvire; Gouze, Jean-Noel; Betz, Oliver B.; Evans, Christopher H.; Ghivizzani, Steven C.
- TI Gene transfer to articular chondrocytes with recombinant adenovirus
- SO Methods in Molecular Biology (Totowa, NJ, United States) (2003), 215 (Cytokines and Colony Stimulating Factors), 235-246 CODEN: MMBIED; ISSN: 1064-3745
- L14 ANSWER 5 OF 36 MEDLINE DUPLICATE 4
- AU Abe Nobuhiro; Lee Yu-Po; Sato Makoto; Zhang Xuguang; Wu Jing; Mitani Kohnosuke; Lieberman Jay R
- TI Enhancement of bone repair with a helper-dependent adenoviral transfer of bone morphogenetic protein-2.
- SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (2002 Sep 27) 297 (3) 523-7.

 Journal code: 0372516. ISSN: 0006-291X.
- L14 ANSWER 6 OF 36 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 5
- AU Jane, John A., Jr.; Dunford, Bradley A.; Kron, Adam; Pittman, Debra D.; Sasaki, Tsutomu; Li, Jin Zhong; Li, Hongwei; Alden, Tord D.; Dayoub, Hayan; Hankins, Gerald R.; Kallmes, David F.; Helm, Gregory A.
- TI Ectopic osteogenesis using adenoviral bone morphogenetic protein (BMP)-4 and BMP-6 gene transfer
- SO Molecular Therapy (2002), 6(4), 464-470 CODEN: MTOHCK; ISSN: 1525-0016
- L14 ANSWER 7 OF 36 MEDLINE DUPLICATE 6
- AU van Griensven M; Lobenhoffer P; Barke A; Tschernig T; Lindenmaier W; Krettek C; Gerich T G
- TI Adenoviral gene transfer in a rat fracture model.
- SO LABORATORY ANIMALS, (2002 Oct) 36 (4) 455-61. Journal code: 0112725. ISSN: 0023-6772.
- L14 ANSWER 8 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 7
- AU Tezuka, Ken-ichi (1); Yasuda, Masafumi; Watanabe, Naoko; Morimura, Naoko; Kuroda, Kazuki; Miyatani, Seiji; Hozumi, Nobumichi
- TI Stimulation of osteoblastic cell differentiation by Notch.
- SO Journal of Bone and Mineral Research, (February, 2002) Vol. 17, No. 2, pp. 231-239. print.
 ISSN: 0884-0431.
- L14 ANSWER 9 OF 36 MEDLINE DUPLICATE 8
- AU Alden Tord D; Varady Peter; Kallmes David F; Jane John A Jr; Helm Gregory
- TI Bone morphogenetic protein gene therapy.
- SO SPINE, (2002 Aug 15) 27 (16 Suppl 1) S87-93. Ref: 102 Journal code: 7610646. ISSN: 1528-1159.
- L14 ANSWER 10 OF 36 CAPLUS COPYRIGHT 2003 ACS
- AU Bondestam, Jonas; Kaivo-Oja, Noora; Kallio, Janne; Groome, Nigel; Hyden-Granskog, Christel; Fujii, Makiko; Moustakas, Aristidis; Jalanko,

- Anu; ten Dijke, Peter; Ritvos, Olli
- TI Engagement of activin and bone morphogenetic protein signaling pathway Smad proteins in the induction of inhibin B production in ovarian granulosa cells
- SO Molecular and Cellular Endocrinology (2002), 195(1-2), 79-88 CODEN: MCEND6; ISSN: 0303-7207
- L14 ANSWER 11 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Tsuru, Michiyo; Nagata, Kensei (1); Ueno, Takato; Jimi, Atsuo; Noda, Shinshi; Iida, Shizuka; Sata, Michio
- TI Confocal laser microscopy of chondrocytes that received gene transfer using in vitro electroporation.
- SO Kurume Medical Journal, (2002) Vol. 49, No. 1-2, pp. 1-5. print. ISSN: 0023-5679.
- L14 ANSWER 12 OF 36 MEDLINE DUPLICATE 9
- AU Gelse K; Jiang Q J; Aigner T; Ritter T; Wagner K; Poschl E; von der Mark K; Schneider H
- TI Fibroblast-mediated delivery of growth factor complementary DNA into mouse joints induces chondrogenesis but avoids the disadvantages of direct viral gene transfer.
- SO ARTHRITIS AND RHEUMATISM, (2001 Aug) 44 (8) 1943-53. Journal code: 0370605. ISSN: 0004-3591.
- L14 ANSWER 13 OF 36 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AU Lindsey W H (Reprint)
- TI Osseous tissue engineering with gene therapy for facial bone reconstruction
- SO LARYNGOSCOPE, (JUL 2001) Vol. 111, No. 7, pp. 1128-1136.
 Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA
 19106-3621 USA.
 ISSN: 0023-852X.
- L14 ANSWER 14 OF 36 MEDLINE DUPLICATE 10
- AU Hidaka C; Quitoriano M; Warren R F; Crystal R G
- TI Enhanced matrix synthesis and in vitro formation of cartilage-like tissue by genetically modified chondrocytes expressing BMP-7.
- SO JOURNAL OF ORTHOPAEDIC RESEARCH, (2001 Sep) 19 (5) 751-8. Journal code: 8404726. ISSN: 0736-0266.
- L14 ANSWER 15 OF 36 MEDLINE DUPLICATE 11
- AU Varady P; Li J Z; Cunningham M; Beres E J; Das S; Engh J; Alden T D; Pittman D D; Kerns K M; Kallmes D F; Helm G A
- TI Morphologic analysis of BMP-9 gene therapy-induced osteogenesis.
- SO HUMAN GENE THERAPY, (2001 Apr 10) 12 (6) 697-710. Journal code: 9008950. ISSN: 1043-0342.
- L14 ANSWER 16 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE
- AU Okubo, Yasunori; Bessho, Kazuhisa (1); Fujimura, Kazuma; Kaihara, Shinji; Iizuka, Tadahiko; Miyatake, Shin-Ichi
- TI The time course study of osteoinduction by bone morphogenetic protein-2 via adenoviral vector.
- SO Life Sciences, (December 7, 2001) Vol. 70, No. 3, pp. 325-336. http://www.elsevier.nl/inca/publications/store/5/2/5/4/7/7/index.htt. print. ISSN: 0024-3205.
- L14 ANSWER 17 OF 36 MEDLINE DUPLICATE 13
- AU Helm G A; Li J Z; Alden T D; Hudson S B; Beres E J; Cunningham M; Mikkelsen M M; Pittman D D; Kerns K M; Kallmes D F
- TI A light and electron microscopic study of ectopic tendon and ligament formation induced by bone morphogenetic protein-13 adenoviral gene therapy.

- SO JOURNAL OF NEUROSURGERY, (2001 Aug) 95 (2) 298-307. Journal code: 0253357. ISSN: 0022-3085.
- L14 ANSWER 18 OF 36 MEDLINE DUPLICATE 14
- AU Turgeman G; Pittman D D; Muller R; Kurkalli B G; Zhou S; Pelled G; Peyser A; Zilberman Y; Moutsatsos I K; Gazit D
- TI Engineered human mesenchymal stem cells: a novel platform for skeletal cell mediated gene therapy.
- SO JOURNAL OF GENE MEDICINE, (2001 May-Jun) 3 (3) 240-51. Journal code: 9815764. ISSN: 1099-498X.
- L14 ANSWER 19 OF 36 MEDLINE DUPLICATE 15
- AU Okubo Y; Bessho K; Fujimura K; Iizuka T; Miyatake S I
- TI In vitro and in vivo studies of a bone morphogenetic protein-2 expressing adenoviral vector.
- SO JOURNAL OF BONE AND JOINT SURGERY. AMERICAN VOLUME, (2001) 83-A Suppl 1 (Pt 2) S99-104.

 Journal code: 0014030. ISSN: 0021-9355.
- L14 ANSWER 20 OF 36 MEDLINE DUPLICATE 16
- AU Cheng S L; Lou J; Wright N M; Lai C F; Avioli L V; Riew K D
- TI In vitro and in vivo induction of bone formation using a recombinant adenoviral vector carrying the human BMP-2 gene.
- SO CALCIFIED TISSUE INTERNATIONAL, (2001 Feb) 68 (2) 87-94.

 Journal code: 7905481. ISSN: 0171-967X.
- L14 ANSWER 21 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 17
- AU Krebsbach, Paul H. (1); Gu, Keni; Franceschi, Renny T.; Rutherford, R. Bruce
- TI Gene therapy-directed osteogenesis: BMP-7-transduced human fibroblasts form bone in vivo.
- SO Human Gene Therapy, (May 20, 2000) Vol. 11, No. 8, pp. 1201-1210. print. ISSN: 1043-0342.
- L14 ANSWER 22 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Smith, P.; Shuler, F. D.; Georgescu, H. I.; Ghivizzani, S. C.; Johnstone, B.; Niyibizi, C.; Robbins, P. D.; Evans, C. H. (1)
- TI Genetic enhancement of matrix synthesis by articular chondrocytes: Comparison of different growth factor genes in the presence and absence of interleukin-1.
- SO Arthritis & Rheumatism, (May, 2000) Vol. 43, No. 5, pp. 1156-1164. print. ISSN: 0004-3591.
- L14 ANSWER 23 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 18
- AU Haberberger, T. C.; Kupfer, K.; Murphy, J. E. (1)
- TI Profiling of genes which are differentially expressed in mouse liver in response to adenoviral vectors and delivered genes.
- SO Gene Therapy, (June, 2000) Vol. 7, No. 11, pp. 903-909. print. ISSN: 0969-7128.
- L14 ANSWER 24 OF 36 CAPLUS COPYRIGHT 2003 ACS
- AU Nakaoka, Takashi; Gonda, Koichi
- TI Gene therapy using BMP-2 gene
- SO Molecular Medicine (Tokyo) (2000), 37(6), 708-714 CODEN: MOLMEL; ISSN: 0918-6557
- L14 ANSWER 25 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 19
- AU Franceschi, Renny T. (1); Wang, Dian; Krebsbach, Paul H.; Rutherford, R. Bruce
- TI Gene therapy for bone formation: In vitro and in vivo osteogenic activity

- of an adenovirus expressing BMP7.
- SO Journal of Cellular Biochemistry, (6 June, 2000) Vol. 78, No. 3, pp. 476-486. print.
 ISSN: 0730-2312.
- L14 ANSWER 26 OF 36 MEDLINE DUPLICATE 20
- AU Okubo Y; Bessho K; Fujimura K; Iizuka T; Miyatake S I
- TI Osteoinduction by bone morphogenetic protein-2 via adenoviral vector under transient immunosuppression.
- SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (2000 Jan 7) 267 (1) 382-7.

 Journal code: 0372516. ISSN: 0006-291X.
- L14 ANSWER 27 OF 36 MEDLINE DUPLICATE 21
- AU Helm G A; Alden T D; Beres E J; Hudson S B; Das S; Engh J A; Pittman D D; Kerns K M; Kallmes D F
- TI Use of bone morphogenetic protein-9 gene therapy to induce spinal arthrodesis in the rodent.
- SO JOURNAL OF NEUROSURGERY, (2000 Apr) 92 (2 Suppl) 191-6. Journal code: 0253357. ISSN: 0022-3085.
- L14 ANSWER 28 OF 36 MEDLINE DUPLICATE 22
- AU Alden T D; Beres E J; Laurent J S; Engh J A; Das S; London S D; Jane J A Jr; Hudson S B; Helm G A
- TI The use of bone morphogenetic protein gene therapy in craniofacial bone repair.
- SO JOURNAL OF CRANIOFACIAL SURGERY, (2000 Jan) 11 (1) 24-30. Journal code: 9010410. ISSN: 1049-2275.
- L14 ANSWER 29 OF 36 CAPLUS COPYRIGHT 2003 ACS
- IN Helm, Gregory A.; Hankins, Gerald R.; Alden, Tord D.; Chung, Leland W. K.; Ko, Song-Chu; Kao, Chinghai
- TI Gene therapy vector with osteocalcin promoter and genes for bone morphogenic proteins or growth factors
- SO PCT Int. Appl., 44 pp. CODEN: PIXXD2
- L14 ANSWER 30 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 23
- AU Piek, Ester; Moustakas, Aristidis (1); Kurisaki, Akira; Heldin, Carl-Henrik; ten Dijke, Peter
- TI TGF-beta type I receptor/ALK-5 and Smad proteins mediate epithelial to mesenchymal transdifferentiation in NMuMG breast epithelial cells.
- SO Journal of Cell Science, (Dec., 1999) Vol. 112, No. 24, pp. 4557-4568. ISSN: 0021-9533.
- L14 ANSWER 31 OF 36 MEDLINE DUPLICATE 24
- AU Alden T D; Pittman D D; Hankins G R; Beres E J; Engh J A; Das S; Hudson S B; Kerns K M; Kallmes D F; Helm G A
- TI In vivo endochondral bone formation using a bone morphogenetic protein 2 adenoviral vector.
- SO HUMAN GENE THERAPY, (1999 Sep 1) 10 (13) 2245-53. Journal code: 9008950. ISSN: 1043-0342.
- L14 ANSWER 32 OF 36 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AU Lieberman J R (Reprint); Daluiski A; Stevenson S; Wu L; McAllister P; Lee Y P; Kabo J M; Finerman G A M; Berk A J; Witte O N
- TI The effect of regional gene therapy with bone morphogenetic protein-2-producing bone-marrow cells on the repair of segmental femoral defects in rats
- SO JOURNAL OF BONE AND JOINT SURGERY-AMERICAN VOLUME, (JUL 1999) Vol. 81A, No. 7, pp. 905-917.
 Publisher: JOURNAL BONE JOINT SURGERY INC, 20 PICKERING ST, NEEDHAM, MA 02192.

ISSN: 0021-9355.

- L14 ANSWER 33 OF 36 MEDLINE DUPLICATE 25
- AU Okubo Y; Bessho K; Fujimura K; Iizuka T; Miyatake S
- TI Expression of bone morphogenetic protein-2 via adenoviral vector in C2C12 myoblasts induces differentiation into the osteoblast lineage.
- SO BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, (1999 Sep 7) 262 (3) 739-43.

 Journal code: 0372516. ISSN: 0006-291X.
- L14 ANSWER 34 OF 36 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 26
- AU Musgrave, D. S.; Bosch, P.; Ghivizzani, S.; Robbins, P. D.; Evans, C. H.; Huard, J. (1)
- TI Adenovirus-mediated direct gene therapy with bone morphogenetic protein-2 produces bone.
- SO Bone (New York), (June, 1999) Vol. 24, No. 6, pp. 541-547. ISSN: 8756-3282.
- L14 ANSWER 35 OF 36 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AU Baltzer A W A (Reprint); Lattermann C; Whalen J D; Braunstein S; Robbins P D; Evans C H
- TI A gene therapy approach to accelerating bone healing Evaluation of gene expression in a New Zealand white rabbit model
- SO KNEE SURGERY SPORTS TRAUMATOLOGY ARTHROSCOPY, (MAY 1999) Vol. 7, No. 3, pp. 197-202.
 Publisher: SPRINGER VERLAG, 175 FIFTH AVE, NEW YORK, NY 10010.
 ISSN: 0942-2056.
- L14 ANSWER 36 OF 36 MEDLINE DUPLICATE 27
- AU Riew K D; Wright N M; Cheng S; Avioli L V; Lou J
- TI Induction of bone formation using a recombinant adenoviral vector carrying the human BMP-2 gene in a rabbit spinal fusion model.
- SO CALCIFIED TISSUE INTERNATIONAL, (1998 Oct) 63 (4) 357-60. Journal code: 7905481. ISSN: 0171-967X.

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(FILE 'HOME' ENTERED AT 15:25:30 ON 28 APR 2003)

FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 15:25:41 ON 28 APR 2003

97859 S WOUND (3A) HEAL? OR EGR

L2 2337 S ADENOVIR? (5A) REPLICATION-DEFICIENT

L3 32 S L1 AND L2

L4 14 DUP REM L3 (18 DUPLICATES REMOVED)

=> d au ti so ab 1-14 14

L4 ANSWER 1 OF 14 CAPLUS COPYRIGHT 2003 ACS

IN Kovesdi, Imre; Kessler, Paul D.

TI Human vascular endothelial growth factor (VEGF) fusion proteins and their therapeutic uses in angiogenesis, bone growth, and wound healing

SO PCT Int. Appl., 191 pp. CODEN: PIXXD2

AB The invention provides therapeutic fusion proteins which include a first peptide portion comprising a first non-heparin binding vascular endothelial growth factor (VEGF) peptide portion and a second non-VEGF peptide portion covalently assocd. with the first peptide portion, which first and second peptide portions sep. promote angiogenesis, bone growth, wound healing, or any combination thereof. Further provided are polynucleotides, encoding such fusion proteins, vectors including such polynucleotides, methods of making such proteins, and methods of promoting angiogenesis, bone growth, and/or wound healing using such proteins, polynucleotides, and vectors. invention specifically claims VEGF fusion proteins with angiopoietin, a fibroblast growth factor, a member of the HBNF-MK family of growth factors, alk. phosphatase, and fusion proteins which lack a collagen binding domain. In addn., the invention claims use of linker peptides to improve fusion protein stability and activities. One example of the invention shows binding of VEGF fusion proteins to the flk-1 VEGF receptor. Another example shows angiogenesis induced by injection of mouse ear or rat hind limb with recombinant adenoviral vectors expressing VEGF fusion proteins. An assay for blood vessel permeability suggests that the angiogenic fusion proteins reduce vascular leakage assocd. with VEGF biol. activity.

- L4 ANSWER 2 OF 14 CAPLUS COPYRIGHT 2003 ACS
- IN Brough, Douglas E.; King, C. Richter; Kovesdi, Imre
- TI Construction of replication deficient adenoviral vectors expressing tumor necrosis factor(TNF) and uses in antitumor therapy
- SO PCT Int. Appl., 45 pp. CODEN: PIXXD2
- The invention discloses methods of constructing replication deficient adenoviral vectors comprising a DNA sequence coding for TNF and their uses in antitumor therapy. In particular, the invention provides a viral vector that comprises an adenoviral genome deficient in the E1, E3 and E4 regions, a DNA sequence coding for TNF, and a radiation inducible promoter operably linked to the DNA sequence coding for TNF. The invention also provides a vector comprising an adenoviral genome deficient in the E1, E3 and E4 regions, a DNA sequence coding for TNF, and a spacer element of at least 15 base pairs in the E4 region of the adenoviral genome. The invention further provides methods of treating a tumor or cancer by administering an effective amt. of the antitumor adenoviral vector.

- AU Romano Di Peppe S; Mangoni A; Zambruno G; Spinetti G; Melillo G; Napolitano M; Capogrossi M C
- Adenovirus-mediated VEGF(165) gene transfer enhances wound healing by promoting angiogenesis in CD1 diabetic mice.
- SO GENE THERAPY, (2002 Oct) 9 (19) 1271-7. Journal code: 9421525. ISSN: 0969-7128.
- AB It has been previously shown that vascular endothelial growth factor (VEGF) plays a central role in promoting angiogenesis during wound repair and that healing-impaired diabetic mice show decreased VEGF expression levels. In order to investigate the potential benefits of gene therapy with growth factors on wound repair, a replication-deficient recombinant adenovirus vector carrying the human VEGF(165) gene (AdCMV.VEGF(165)) was topically applied on excisional wounds of streptozotocin-induced diabetic mice. Treatment with AdCMV.VEGF(165) significantly accelerated wound closure when compared with AdCMV.LacZ-treated, as well as saline-treated control mice, by promoting angiogenesis at the site of injury. Our findings suggest that AdCMV.VEGF(165) may be regarded as a therapeutic tool for the treatment of diabetic ulcers.
- L4 ANSWER 4 OF 14 MEDLINE

- AU Perkins Todd W; Faha Barbara; Ni Ming; Kiland Julie A; Poulsen Gretchen L; Antelman Doug; Atencio Isabella; Shinoda Jeremy; Sinha Dinesh; Brumback Lyndia; Maneval Daniel; Kaufman Paul L; Nickells Robert W
- Adenovirus-mediated gene therapy using human p21WAF-1/Cip-1 to prevent wound healing in a rabbit model of glaucoma filtration surgery.
- SO ARCHIVES OF OPHTHALMOLOGY, (2002 Jul) 120 (7) 941-9. Journal code: 7706534. ISSN: 0003-9950.
- OBJECTIVE: To determine if adenovirus-mediated p21(WAF-1/Cip-1) (p21) gene AB therapy can prevent fibroproliferation and wound healing in a rabbit model of glaucoma filtration surgery. METHODS: In vitro studies were performed using rabbit Tenon fibroblasts harvested from fresh tissue. In vivo studies were conducted in New Zealand white rabbits. A full-thickness sclerotomy was performed under a limbal-based conjunctival flap. Reagents tested included a replication-deficient recombinant adenovirus containing the human p21 gene (rAd.p21); the nonspecific marker gene for green fluorescent protein or beta-galactosidase; mitomycin, 0.5 mg/mL; and balanced saline solution. Each treatment was applied episclerally for 5 minutes before the sclerotomy using a soaked cellulose sponge placed under the surgically created conjunctival flap. Independent experiments were conducted to (1) monitor changes in intraocular pressure during a 30-day period after treatment and examine surgical site histological features, (2) examine changes in bleb morphologic features over 30 days, (3) determine outflow facility 14 days after treatment, and (4) examine the localization and persistence of rAd.p21 expression between 3 and 60 days after treatment. RESULTS: Treatment of Tenon fibroblasts with rAd.p21 resulted in a dose-dependent inhibition of DNA synthesis and cell growth in vitro. vivo, rAd.p21 inhibited wound healing and fibroproliferation after filtration surgery, comparably to mitomycin. Mitomycin caused notable thinning of the bleb wall. In addition, 2 of the 5 mitomycin-treated eyes exhibited an abscess with hypopyon and hyalitis 30 days after surgery, which was not observed in any of the rAd.p21-treated eyes. None of the treatments resulted in a significantly sustained decrease in intraocular pressure during the 30-day period, although mitomycin treatment resulted in a significant (P = .02) increase in outflow facility 2 weeks after surgery in separate animals. Mitomycinand rAd.p21-treated eyes had functioning blebs at the end of the experiment based on slitlamp examination. CONCLUSIONS: Mitomycin and rAd.p21 were effective in preventing fibroproliferation and wound healing in a rabbit model of glaucoma surgery. Mitomycin treatment increased outflow facility in normal-pressure eyes. RELEVANCE: Gene therapy with rAd.p21 may provide an effective

antiproliferative for glaucoma filtration surgery, without the complications associated with mitomycin.

L4 ANSWER 5 OF 14 MEDLINE

- DUPLICATE 3
- AU Gupta Vinay K; Park James O; Jaskowiak Nora T; Mauceri Helena J; Seetharam Saraswathy; Weichselbaum Ralph R; Posner Mitchell C
- TI Combined gene therapy and ionizing radiation is a novel approach to treat human esophageal adenocarcinoma.
- SO ANNALS OF SURGICAL ONCOLOGY, (2002 Jun) 9 (5) 500-4. Journal code: 9420840. ISSN: 1068-9265.
- BACKGROUND: The ability to infect tumor cells limits the antitumor effects AB of gene therapy. The addition of radiotherapy to treatment with Ad. Egr. TNF. 11D, a replication-deficient adenovirus containing a radiation-inducible promoter, early growth response-1, and the tumor necrosis factor-alpha (TNFalpha) complementary DNA may enhance the therapeutic ratio. METHODS: Seg-1 human esophageal adenocarcinoma cells were treated with Ad. Egr. TNF. 11D with or without radiation. TNFalpha levels were quantified with enzyme-linked immunosorbent assay. Athymic nude mice bearing Seg-1 tumors were randomized to buffer, ionizing radiation, Ad. Egr. TNF. 11D, and combination therapy. Tumor growth delay was used to compare treatment regimens. TNFalpha levels were measured in tumor homogenates and plasma. RESULTS: Seg-1 cells treated with Ad. Egr. TNF. 11D and ionizing radiation demonstrated increased TNFalpha levels at 72 hours compared with cells exposed to vector alone (124 +/- 0 pg/mL vs. 31.11 +/- 22 pg/mL; P $^{\circ}$ =.008). In vivo, Ad. Egr. TNF.11D-treated tumors expressed low TNFalpha levels (151.5 +/- 107.11 pg/mg protein) compared with tumors receiving combined treatment (793.92 \pm 489.13 pg/mg protein; P = .067). Increased TNFalpha levels were associated with increased tumor growth delay after combined treatment (P <.05). CONCLUSIONS: Radiotherapy enables focal stimulation of TNFalpha expression in Ad. Egr .TNF.11D-infected cells and thus improves local tumor control.
- L4 ANSWER 6 OF 14 MEDLINE

DUPLICATE 4

- AU Park James O; Lopez Carlos A; Gupta Vinay K; Brown Charles K; Mauceri Helena J; Darga Thomas E; Manan Abdullah; Hellman Samuel; Posner Mitchell C; Kufe Donald W; Weichselbaum Ralph R
- TI Transcriptional control of viral gene therapy by cisplatin.
- SO JOURNAL OF CLINICAL INVESTIGATION, (2002 Aug) 110 (3) 403-10. Journal code: 7802877. ISSN: 0021-9738.
- Ionizing radiation (IR) and radical oxygen intermediates (ROIs) activate AB the early growth response-1 (Egr1) promoter through specific cis-acting sequences termed CArG elements. Ad. Egr. TNF. 11D, a replication-deficient adenoviral vector containing CArG elements cloned upstream of the cDNA for human recombinant TNF-alpha was used to treat human esophageal adenocarcinoma and rat colon adenocarcinoma cells in culture and as xenografts in athymic nude mice. Cisplatin, a commonly used chemotherapeutic agent, causes tumor cell death by producing DNA damage and generating ROIs. The present studies demonstrate induction of TNF-alpha production in tumor cells and xenografts treated with the combination of Ad. Egr. TNF.11D and cisplatin. The results show that the Egr1 promoter is induced by cisplatin and that this induction is mediated in part through the CArG elements. These studies also demonstrate an enhanced antitumor response without an increase in toxicity following treatment with Ad. Egr .TNF.11D and cisplatin, compared with either agent alone. Chemo-inducible cancer gene therapy thus provides a means to control transgene expression while enhancing the effectiveness of commonly used chemotherapeutic
- L4 ANSWER 7 OF 14 MEDLINE

agents.

DUPLICATE 5

AU Danjo Yukitaka; Gipson Ilene K

TI Specific transduction of the leading edge cells of migrating epithelia demonstrates that they are replaced during healing.

- SO EXPERIMENTAL EYE RESEARCH, (2002 Feb) 74 (2) 199-204. Journal code: 0370707. ISSN: 0014-4835.
- As wounds in stratified epithelia close, the numbers of cells at the leading edge of migration decreases. It is not known whether cells at the leading edge are continually replaced or whether some retain their position at the leading edge over time. Replication-deficient adenovirus carrying the green fluorescent protein gene was applied to corneal epithelial wounds in mice and it was found that they primarily infect the leading edge cells of healing epithelium. Eighteen hr after viral transduction, green fluorescent protein expressing cells were located in the apical layer at varying distances behind the leading edge. These data indicate that leading edge cells are replaced during healing of stratified epithelia. Copyright 2002 Elsevier Science Ltd.
- L4 ANSWER 8 OF 14 MEDLINE

- AU Spector J A; Mehrara B J; Luchs J S; Greenwald J A; Fagenholz P J; Saadeh P B; Steinbrech D S; Longaker M T
- TI Expression of adenovirally delivered gene products in healing osseous tissues.
- SO ANNALS OF PLASTIC SURGERY, (2000 May) 44 (5) 522-8.
 Journal code: 7805336. ISSN: 0148-7043.
- Gene therapy has moved from the promise of laboratory investigation to the AB reality of clinical practice in just the last decade. Various methods for delivery of genes to host cells have been developed and utilized both in vitro and in vivo. From the perspective of the plastic surgeon, gene therapy holds the promise to augment healing in clinical situations that remain difficult to treat, such as chronic wounds, osteoradionecrosis, or possibly to expedite current clinical practices, such as distraction osteogenesis. The authors chose to investigate the potential for gene therapy in osseous tissues using a replication-deficient adenovirus vector to deliver the marker transgene beta-galactosidase. An adenovirus vector is ideal for use in situations in which transgene expression is desired for only a relatively short period of time, such as wound and fracture healing. Utilizing a rat mandibular osteotomy model, they demonstrated that, using an adenoviral vector, foreign genes can be delivered in a simple fashion and can be expressed in a reliable manner within and around the osteotomy site for at least 10 days. Furthermore, there was no evidence of transfection of distant tissues associated with local application of the adenovirus vector. With this information, clinicians may now attempt to deliver osteogenic and angiogenic genes in a site-specific fashion to improve and expedite osseous healing.
- L4 ANSWER 9 OF 14 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Manome, Yoshinobu (1); Kunieda, Takehiko; Wen, Patrick Y.; Koga, Tomohiko; Kufe, Donald W.; Ohno, Tsuneya
- TI Transgenic expression in malignant glioma using a replication-defective adenoviral vector containing the **Egr**-1 promoter: Activation by ionizing radiation or uptake of radioactive iododeoxyuridine.
- SO Human Gene Therapy, (July 1, 1998) Vol. 9, No. 10, pp. 1409-1417. ISSN: 1043-0342.
- One approach to improving the specificity of gene therapy involves using radiosensitive promoters to activate gene expression selectively in the radiation field. In this study, we evaluated the ability of irradiation to regulate the transcription of a recombinant replication-defective adenovirus vector, Ad.Egr-1/lacZ, containing the radiation-inducible Egr-1 promoter driving the beta-galactosidase reporter gene in glioma cells. Transcripts of the Egr-1 gene in human and rat glioma cells were induced following irradiation with as little as 2 Gy. This dose was 10-fold less than previously reported, and comparable to doses of irradiation used clinically in standard fractionated radiotherapy for brain tumors. When 9L rat gliosarcoma cells were infected with Ad.Egr1/lacZ in vitro and exposed

to 2 Gy of external beam irradiation, there was a threefold increase in beta-galactosidase expression. Irradiation of intracerebral 9L tumors infected with the Ad.Egr-1/lacZ virus, using either external beam radiotherapy (2 Gy) or the thymidine analog 5-iodo-2'-deoxyuridine radiolabeled with the Auger electron emitter iodine-125 ((125I)IdUrd), also resulted in increased beta-galactosidase activity of the tumor cells. These results indicate that the use of viral vectors containing radiation-inducible promoters represents a novel therapeutic approach that enables gene therapy to be spatially and temporally regulated by ionizing radiation. These findings also support a potential role for radiation-inducible promoters in the treatment of malignant brain tumors.

- L4 ANSWER 10 OF 14 CAPLUS COPYRIGHT 2003 ACS
- AU Kitano, Yukie; Radu, Antoneta; Sylvester, Karl G.; Nesbit, Mark; Herlyn, Meenhard; Adzick, N. Scott; Crombleholme, Timothy M.
- TI Treatment of large excisional healing-impaired wounds in diabetic mice with a replication-deficient adenovirus carrying the PDGF-B transgene (PDGF-B/Adv5-CMV)
- SO Surgical Forum (1998), 49, 651-653 CODEN: SUFOAX; ISSN: 0071-8041
- AB The authors hypothesized that the induction of PDGF-B gene over-expression in cells participating in wound healing by adenoviral-mediated gene transfer of human PDGF-B transgene would accelerate wound healing. The results showed that adenoviral-mediated gene transfer of PDGF-B may be a more effective growth factor administration in healing impaired wounds in diabetics.
- L4 ANSWER 11 OF 14 MEDLINE DUPLICATE 7
- AU Chung T D; Mauceri H J; Hallahan D E; Yu J J; Chung S; Grdina W L; Yajnik S; Kufe D W; Weichselbaum R R
- TI Tumor necrosis factor-alpha-based gene therapy enhances radiation cytotoxicity in human prostate cancer.
- SO CANCER GENE THERAPY, (1998 Nov-Dec) 5 (6) 344-9. Journal code: 9432230. ISSN: 0929-1903.
- AB The purpose of the present study was to determine the therapeutic potential of combining radiotherapy with tumor necrosis factor (TNF)-alpha-based gene therapy in the human prostate cancer PC-3 xenograft. PC-3 cells are highly resistant to TNF-alpha-induced cytotoxicity in vitro. A modest enhancement of radiation killing was observed with the addition of TNF-alpha in clonogenic survival assays. Combined treatment with Ad. Egr-TNF, a replicationdeficient adenovirus modified to express TNF-alpha following the exposure of infected cells to ionizing radiation (40 Gy administered at 5 Gy per fraction) in vivo, resulted in increased tumor control, as defined by a reduction of tumor volume, when compared with treatment with Ad.Egr-TNF alone or with radiation alone (P < .03). The improvement in tumor control was achieved without increasing acute normal tissue damage when compared with tissue injury from radiation alone. The results of these studies support further development and clinical application of genetic radiotherapy for human prostate cancer.
- L4 ANSWER 12 OF 14 MEDLINE
- AU Lou J; Kubota H; Hotokezaka S; Ludwig F J; Manske P R
- TI In vivo gene transfer and overexpression of focal adhesion kinase (pp125 FAK) mediated by recombinant adenovirus-induced tendon adhesion formation and epitenon cell change.
- SO JOURNAL OF ORTHOPAEDIC RESEARCH, (1997 Nov) 15 (6) 911-8. Journal code: 8404726. ISSN: 0736-0266.
- AB Adhesion formation is a frequent complication of tendon injury repair: however, little is known about its mechanisms. The intracellular focal adhesion kinase (FAK)-related signaling pathway may be one of the mechanisms involved in the induction of tendon adhesions. The replication deficient adenovirus containing

the FAK gene (pp125 FAK) was constructed and named Adv-Fak. By in vitro transductions with the recombinant virus, overexpression of the FAK protein was documented in transduced cultured primary tendon cells. By in vivo direct injection of Adv-FAK into the space between the tendon and tendon sheath of White Leghorn chickens, FAK gene transfer with overexpression of the FAK protein was detected by immunohistological staining. The morphology of these stained cells changed from the normal flat shape to cuboid. The group with overexpressed adenovirus-mediated FAK had significant adhesion formation, as seen by increased work of flexion (118.197 +/- 29.616), compared with the group with overexpressed adenovirus-mediated beta-galactosidase (67.507 \pm - 36.066) (p < 0.0393) and the group with adenovirus-mediated FAK antisense gene transfer (60.357 +/-48.562) (p < 0.0211). Histological examination of the samples from tendons with Adv-FAK showed fibers between the tendon and tendon sheath; there were no fibers in the cavities of samples of injured tendons infected with Adv-beta gal. Moreover, at the application site of the former tendons, a thick fiber layer without epitenon cells was built up on the outer surface, whereas a thin fiber layer with clear epitenon cells was observed in the tendons to which Adv-beta gal was applied. Our results show that overexpression of FAK can induce tendon adhesion formation in vivo. This indicates that FAK and the FAK-related signaling pathway may be involved in the process of tendon adhesion formation. Understanding the details of this process may help to prevent tendon adhesion and improve healing.

- L4 ANSWER 13 OF 14 CAPLUS COPYRIGHT 2003 ACS
- IN Finkel, Toren; Epstein, Stephen E.; Crystal, Ronald G.; Guzman, Raul J.
- TI Selective adenoviral mediated gene transfer into vascular neointima
- SO PCT Int. Appl., 48 pp. CODEN: PIXXD2
- Disclosed is a method of selectively expressing DNA in neointimal cells in an injured blood vessel of a subject. The method comprises administering a replication-deficient recombinant adenovirus which delivers the DNA to the blood vessel at the site of injury, such that the adenovirus remains at the site of injury for a time sufficient for the adenovirus to selectively infect neointimal cells and thereby selectively express the DNA in neointimal cells. The DNA encoding a protein or an antisense RNA can be delivered. This method can be used to treat restenosis and, relatedly, to prevent neointimal cell proliferation.
- L4 ANSWER 14 OF 14 MEDLINE DUPLICATE 8
- AU Hallahan D E; Mauceri H J; Seung L P; Dunphy E J; Wayne J D; Hanna N N; Toledano A; Hellman S; Kufe D W; Weichselbaum R R
- TI Spatial and temporal control of gene therapy using ionizing radiation.
- SO NATURE MEDICINE, (1995 Aug) 1 (8) 786-91. Journal code: 9502015. ISSN: 1078-8956.
- Activation of transcription of the Egr-1 gene by X-rays is AB regulated by the promoter region of this gene. We linked the radiation-inducible promoter region of the Egr-1 gene to the gene encoding the radiosensitizing and tumoricidal cytokine, tumour necrosis factor-alpha (TNF-alpha) and used a replicationdeficient adenovirus to deliver the Egr-TNF construct to human tumours growing in nude mice. Combined treatment with Ad5.Egr-TNF and 5,000 cGy (rad) resulted in increased intratumoral TNF-alpha production and increased tumour control compared with treatment with Ad5.Egr-TNF alone or with radiation alone. The increase in tumour control was achieved without an increase in normal tissue damage when compared to tissue injury from radiation alone. Control of gene transcription by ionizing radiation in vivo represents a novel method of spatial and temporal regulation of gene-based medical treatments.

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L5
          17729 ADENOVIR? (W) VECTOR
 => s l1 and l5
L6
           137 L1 AND L5
=> dup rem 16
PROCESSING COMPLETED FOR L6
              84 DUP REM L6 (53 DUPLICATES REMOVED)
=> d his
      (FILE 'HOME' ENTERED AT 15:25:30 ON 28 APR 2003)
     FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 15:25:41 ON 28 APR
     2003
L1
          97859 S WOUND (3A) HEAL? OR EGR
           2337 S ADENOVIR? (5A) REPLICATION-DEFICIENT
L2
              32 S L1 AND L2
L3
             14 DUP REM L3 (18 DUPLICATES REMOVED)
L4
          17729 S ADENOVIR? (W) VECTOR
L_5
1.6
            137 S L1 AND L5
L7
              84 DUP REM L6 (53 DUPLICATES REMOVED)
=> s 11(s)15
            75 L1(S) L5
=> dup rem 18
PROCESSING COMPLETED FOR L8
L9
             37 DUP REM L8 (38 DUPLICATES REMOVED)
=> d his
     (FILE 'HOME' ENTERED AT 15:25:30 ON 28 APR 2003)
     FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 15:25:41 ON 28 APR
     2003
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           2337 S ADENOVIR? (5A) REPLICATION-DEFICIENT
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L4
             14 DUP REM L3 (18 DUPLICATES REMOVED)
L_5
          17729 S ADENOVIR? (W) VECTOR
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            137 S L1 AND L5
L7
             84 DUP REM L6 (53 DUPLICATES REMOVED)
L8
             75 S L1(S)L5
L9
             37 DUP REM L8 (38 DUPLICATES REMOVED)
=> d au ti so 1-37 19
L9
     ANSWER 1 OF 37
                        MEDLINE
                                                         DUPLICATE 1
ΑU
     Gruss Claus J; Satyamoorthy Kapaettu; Berking Carola; Lininger John;
     Nesbit Mark; Schaider Helmut; Liu Zhao-June; Oka Masahiro; Hsu Mei-Yu;
     Shirakawa Takashi; Li Gang; Bogenrieder Thomas; Carmeliet Peter; El-Deiry
     Wafik S; Eck Stephen L; Rao Justi S; Baker Andrew H; Bennet Jean T;
     Crombleholme Timothy M; Velazquez Omaida; Karmacharya Jagajan; Margolis
     David J; Wilson James M; Detmar Michael; Skobe Mihaela; Robbins Paul D;
     Buck Clayton; Herlyn Meenhard
TI
     Stroma formation and angiogenesis by overexpression of growth factors,
     cytokines, and proteolytic enzymes in human skin grafted to SCID mice.
     JOURNAL OF INVESTIGATIVE DERMATOLOGY, (2003 Apr) 120 (4) 683-92.
SO
     Journal code: 0426720. ISSN: 0022-202X.
L9
     ANSWER 2 OF 37 CAPLUS COPYRIGHT 2003 ACS
     Kovesdi, Imre; Kessler, Paul D.
IN
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Human vascular endothelial growth factor (VEGF) fusion proteins and their

TI

- therapeutic uses in angiogenesis, bone growth, and wound healing
- SO PCT Int. Appl., 191 pp. CODEN: PIXXD2
- L9 ANSWER 3 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Weichselbaum, Ralph R.; Kufe, Donald W.; Gupta, Vinay; Mauceri, Helen;
 Park, James; Posner, Mitchell
- TI Chemotherapeutic induction of Egr-1 promoter activity with anticancer agents for the treatment of hyperproliferative diseases and cancers
- SO PCT Int. Appl., 84 pp. CODEN: PIXXD2
- L9 ANSWER 4 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Tennenbaum, Tamar; Sampson, Sanford; Kuroki, Toshio; Alt, Addy; Shen, Shlomzion
- TI Protein kinase C and methods and compositions for healing wounds
- SO PCT Int. Appl., 93 pp. CODEN: PIXXD2
- L9 ANSWER 5 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Brough, Douglas E.; King, C. Richter; Kovesdi, Imre
- TI Construction of replication deficient adenoviral vectors expressing tumor necrosis factor(TNF) and uses in antitumor therapy
- SO PCT Int. Appl., 45 pp. CODEN: PIXXD2
- L9 ANSWER 6 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 2
- AU Di Peppe, S. Romano; Mangoni, A.; Zambruno, G.; Spinetti, G.; Melillo, G.; Napolitano, M.; Capogrossi, M. C. (1)
- TI Adenovirus-mediated VEGF165 gene transfer enhances wound healing by promoting angiogenesis in CD1 diabetic mice.
- SO Gene Therapy, (October, 2002) Vol. 9, No. 19, pp. 1271-1277. http://www.naturesj.com/gt/. print. ISSN: 0969-7128.
- L9 ANSWER 7 OF 37 MEDLINE

DUPLICATE 3

- AU Weichselbaum Ralph R; Kufe Donald W; Hellman Samuel; Rasmussen Henrik S; King C Richter; Fischer Paul H; Mauceri Helena J
- TI Radiation-induced tumour necrosis factor-alpha expression: clinical application of transcriptional and physical targeting of gene therapy.
- SO Lancet Oncol, (2002 Nov) 3 (11) 665-71. Ref: 75 Journal code: 100957246. ISSN: 1470-2045.
- L9 ANSWER 8 OF 37 MEDLINE

DUPLICATE 4

- AU Mauceri Helena J; Seetharam Saraswathy; Beckett Michael A; Lee John Y; Gupta Vinay K; Gately Stephen; Stack M Sharon; Brown Charles K; Swedberg Kirsten; Kufe Donald W; Weichselbaum Ralph R
- TI Tumor production of angiostatin is enhanced after exposure to TNF-alpha.
- SO INTERNATIONAL JOURNAL OF CANCER, (2002 Feb. 1) 97 (4) 410-5. Journal code: 0042124. ISSN: 0020-7136.
- L9 ANSWER 9 OF 37 MEDLINE

- AU Park James O; Lopez Carlos A; Gupta Vinay K; Brown Charles K; Mauceri Helena J; Darga Thomas E; Manan Abdullah; Hellman Samuel; Posner Mitchell C; Kufe Donald W; Weichselbaum Ralph R
- TI Transcriptional control of viral gene therapy by cisplatin.
- SO JOURNAL OF CLINICAL INVESTIGATION, (2002 Aug) 110 (3) 403-10. Journal code: 7802877. ISSN: 0021-9738.
- L9 ANSWER 10 OF 37 MEDLINE DUPLICATE 6
- AU Rivera Arnold D C; Walker Charles N; Bleustein Clifford; Choi Benjamin; Poppas Dix P; Felsen Diane
- TI Enhanced adenoviral-vector mediated gene transfer using human albumin

solder.

- SO LASERS IN SURGERY AND MEDICINE, (2002) 30 (4) 313-9. Journal code: 8007168. ISSN: 0196-8092.
- L9 ANSWER 11 OF 37 MEDLINE DUPLICATE 7
- AU Ailawadi Maneesh; Lee Jay M; Lee Sang; Hackett Neil; Crystal Ronald G; Korst Robert J
- TI Adenovirus vector-mediated transfer of the vascular endothelial growth factor cDNA to healing abdominal fascia enhances vascularity and bursting strength in mice with normal and impaired wound healing.
- SO SURGERY, (2002 Feb) 131 (2) 219-27. Journal code: 0417347. ISSN: 0039-6060.
- L9 ANSWER 12 OF 37 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AU Chin G; Gowda S; Schultz G (Reprint)
- TI Evaluation of platelet-derived growth factor in a rat model of ischemic skin wound healing
- SO WOUNDS-A COMPENDIUM OF CLINICAL RESEARCH AND PRACTICE, (JUN 2002) Vol. 14, No. 5, pp. 199-203.

 Publisher: H M P COMMUNICATIONS, 83 GENERAL WARREN BLVD, STE 100, MALVERN, PA 19355 USA.
 ISSN: 1044-7946.
- L9 ANSWER 13 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Gao, C. Y. (1); Negash, S. (1); Zelenka, P. S. (1)
- TI Cdk5 Regulates Corneal Epithelial Cell Adhesion and Migration in vitro.
- SO ARVO Annual Meeting Abstract Search and Program Planner, (2002) Vol. 2002, pp. Abstract No. 4211. cd-rom.

 Meeting Info.: Annual Meeting of the Association For Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA May 05-10, 2002
- L9 ANSWER 14 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Chen, Z. Z. (1); Wen, S.; Maneval, D. (1); Hess, M. (1); Nery, J.; Kaufman, P.; Nickells, R.; Faha, B. (1)
- TI Biodistribution of an Adenovirus Encoding Human p21WAF1/Cip-1(rAd-p21) Following Subconjunctival Injection in Rabbits.
- SO ARVO Annual Meeting Abstract Search and Program Planner, (2002) Vol. 2002, pp. Abstract No. 3334. cd-rom.

 Meeting Info.: Annual Meeting of the Association For Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA May 05-10, 2002
- L9 ANSWER 15 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Havenga, Menzo Jans Emco; Bout, Abraham
- TI Methods and means for enhancing skin transplantation using gene delivery vehicles having tropism for primary fibroblasts, as well as other uses thereof
- SO Eur. Pat. Appl., 26 pp. CODEN: EPXXDW
- L9 ANSWER 16 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 8
- AU Honma, Kimi; Ochiya, Takahiro; Nagahara, Shunji; Sano, Akihiko; Yamamoto, Hanako; Hirai, Kotaro; Aso, Yu; Terada, Masaaki (1)
- TI Atelocollagen-based gene transfer in cells allows high-throughput screening of gene functions.
- SO Biochemical and Biophysical Research Communications, (December 21, 2001) Vol. 289, No. 5, pp. 1075-1081. http://www.academicpress.com/bbrc. print. ISSN: 0006-291X.
- L9 ANSWER 17 OF 37 CAPLUS COPYRIGHT 2003 ACS
- AU Wei, Daoyan; Dai, Bingbing; Chen, Shishu
- TI Tumor targeted expression of adenovirus-mediated CDglyTK gene regulated by irradiation via Egr-1 promoter

- SO Zhonghua Yixue Zazhi (Beijing, China) (2001), 81(16), 999-1003 CODEN: CHHTAT; ISSN: 0376-2491
- L9 ANSWER 18 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 9
- AU Varda-Bloom, N.; Shaish, A.; Gonen, A.; Levanon, K.; Greenbereger, S.; Ferber, S.; Levkovitz, H.; Castel, D.; Goldberg, I.; Afek, A.; Kopolovitc, Y.; Harats, D. (1)
- TI Tissue-specific gene therapy directed to tumor angiogenesis.
- SO Gene Therapy, (June, 2001) Vol. 8, No. 11, pp. 819-827. print. ISSN: 0969-7128.
- L9 ANSWER 19 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 10
- AU Rauma, Tanja; Kumpumaki, Sanna; Anderson, Richard; Davidson, Beverly L.; Ruotsalainen, Heli; Myllyla, Raili; Hautala, Timo (1)
- TI Adenoviral gene transfer restores lysyl hydroxylase activity in type VI Ehlers-Danlos syndrome.
- SO Journal of Investigative Dermatology, (April, 2001) Vol. 116, No. 4, pp. 602-605. print.
 ISSN: 0022-202X.
- L9 ANSWER 20 OF 37 MEDLINE DUPLICATE 11
- AU Hallahan D E; Qu S; Geng L; Cmelak A; Chakravarthy A; Martin W; Scarfone C; Giorgio T
- TI Radiation-mediated control of drug delivery.
- SO AMERICAN JOURNAL OF CLINICAL ONCOLOGY, (2001 Oct) 24 (5) 473-80. Ref: 62 Journal code: 8207754. ISSN: 0277-3732.
- L9 ANSWER 21 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Wuerschmidt, F. (1); Gomaa, I. E. O. (1); Anton, M.; Lukowicz, T. V. (1); Molls, M. (1); Gaensbacher, B.
- TI Gene expression under control of the radiation-inducible Egr-1 promoter in an adenoviral vector: Vector optimization for reduction of unspecific gene expression in the absence of irradiation.
- SO European Journal of Cancer, (October, 2001) Vol. 37, No. Supplement 6, pp. S160. http://www.elsevier.com/locate/ejca. print.

 Meeting Info.: 11th European Cancer Conference Lisbon, Portugal October 21-25, 2001
 ISSN: 0959-8049.
- L9 ANSWER 22 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AU Mauceri, H. J. (1); Seetharam, S.; Beckett, M. A.; Lee, J. Y.; Gately, S.; Stack, M. S.; Gupta, V. K.; Kufe, D. W.; Weichselbaum, R. R.
- TI Treatment with the adenoviral vector Ad. Egr -TNF-alpha enhances plasma angiostatin levels.
- SO Proceedings of the American Association for Cancer Research Annual Meeting, (March, 2001) Vol. 42, pp. 32. print.

 Meeting Info.: 92nd Annual Meeting of the American Association for Cancer Research New Orleans, LA, USA March 24-28, 2001
 ISSN: 0197-016X.
- L9 ANSWER 23 OF 37 MEDLINE DUPLICATE 12
- AU Liu W; Mehrara B J; Chin G S; Hsu M; Peled Z; Longaker M T
- TI The use of newborn rats and an adenoviral gene delivery vector as a model system for wound-healing research.
- SO ANNALS OF PLASTIC SURGERY, (2000 May) 44 (5) 543-51; discussion 551-2. Journal code: 7805336. ISSN: 0148-7043.
- L9 ANSWER 24 OF 37 MEDLINE DUPLICATE 13
- AU Spector J A; Mehrara B J; Luchs J S; Greenwald J A; Fagenholz P J; Saadeh P B; Steinbrech D S; Longaker M T
- TI Expression of adenovirally delivered gene products in healing osseous tissues.

- SO ANNALS OF PLASTIC SURGERY, (2000 May) 44 (5) 522-8.
 Journal code: 7805336. ISSN: 0148-7043.
- L9 ANSWER 25 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 14
- AU Radfar, Ali J.; Robbins, Paul D.; Huard, Johnny; Rosas, Fabio R.; Dohar, Joseph E.; Hebda, Patricia A. (1)
- TI Transplantation of virally transduced cells into the dermis of immunocompetent and immunodeficient (SCID) mice to determine gene expression profile and differential donor cell survival.
- SO Wound Repair and Regeneration, (November December, 2000) Vol. 8, No. 6, pp. 503-510. print.
 ISSN: 1067-1927.
- L9 ANSWER 26 OF 37 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AU Chandler L A (Reprint); Doukas J; Gonzalez A M; Hoganson D K; Gu D L; Ma C L; Nesbit M; Crombleholme T M; Herlyn M; Sosnowski B A; Pierce G F
- TI FGF2-targeted adenovirus encoding platelet-derived growth factor-B enhances de Novo tissue formation
- SO MOLECULAR THERAPY, (AUG 2000) Vol. 2, No. 2, pp. 153-160. Publisher: ACADEMIC PRESS INC, 525 B ST, STE 1900, SAN DIEGO, CA 92101-4495. ISSN: 1525-0016.
- L9 ANSWER 27 OF 37 MEDLINE DUPLICATE 15
- AU Sylvester K G; Nesbit M; Radu A; Herlyn M; Adzick N S; Crombleholme T M
- TI Adenoviral-mediated gene transfer in wound healing: acute inflammatory response in human skin in the SCID mouse model.
- SO WOUND REPAIR AND REGENERATION, (2000 Jan-Feb) 8 (1) 36-44. Journal code: 9310939. ISSN: 1067-1927.
- L9 ANSWER 28 OF 37 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 16
- AU Liechty, Kenneth W.; Sablich, Timothy J.; Adzick, N. Scott; Crombleholme, Timothy M. (1)
- TI Recombinant adenoviral mediated gene transfer in ischemic impaired wound healing.
- SO Wound Repair and Regeneration, (May-June, 1999) Vol. 7, No. 3, pp. 148-153.
 ISSN: 1067-1927.
- L9 ANSWER 29 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Bohlen, Peter
- TI Truncated vascular endothelial cell growth factor-related proteins, VRP-encoding adenoviral vectors, and pharmaceutical use of VRPs
- SO PCT Int. Appl., 113 pp. CODEN: PIXXD2
- L9 ANSWER 30 OF 37 CAPLUS COPYRIGHT 2003 ACS
- IN Billiar, Timothy R.; Tzeng, Edith; Shears, Larry L., II; Geller, David A.;
 Edington, Howard David James
- TI Methods for promoting wound healing and treating transplant-associated vasculopathy
- SO PCT Int. Appl., 55 pp. CODEN: PIXXD2
- L9 ANSWER 31 OF 37 MEDLINE DUPLICATE 17
- AU Manome Y; Kunieda T; Wen P Y; Koga T; Kufe D W; Ohno T
- TI Transgene expression in malignant glioma using a replication-defective adenoviral vector containing the Egr-1 promoter: activation by ionizing radiation or uptake of radioactive iododeoxyuridine.
- SO HUMAN GENE THERAPY, (1998 Jul 1) 9 (10) 1409-17. Journal code: 9008950. ISSN: 1043-0342.

- L9 ANSWER 32 OF 37 MEDLINE DUPLICATE 18
- AU Yamasaki K; Edington H D; McClosky C; Tzeng E; Lizonova A; Kovesdi I; Steed D L; Billiar T R
- TI Reversal of impaired wound repair in iNOS-deficient mice by topical adenoviral-mediated iNOS gene transfer.
- SO JOURNAL OF CLINICAL INVESTIGATION, (1998 Mar 1) 101 (5) 967-71. Journal code: 7802877. ISSN: 0021-9738.
- L9 ANSWER 33 OF 37 MEDLINE DUPLICATE 19
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